CLAIM AMENDMENTS

1-13. (Cancelled)

- 14. (Currently amended) A pharmaceutical Pharmaceutical composition comprising (a) a therapeutically effective amount of an active principle for oral use that is in the form of a system which is and (b) a self micro-emulsifying carrier, said self micro-emulsifying carrier self-micro-emulsifying on contact with an aqueous phase, comprising:
 - a therapeutically effective amount of the said active principle;
- (i) [[-]] a lipophilic phase comprising a mixture of glycerol mono-, di- and triesters and of PEG mono- and diesters with at least one fatty acid chosen from the group comprising consisting of C₈-C₁₈ fatty acids;
- (ii) [[-]] a surfactant phase comprising a mixture of glycerol mono-, di- and triesters and of PEG mono- and diesters with caprylic acid (C_8) and capric acid (C_{10});
- (iii) a co-surfactant phase comprising at least one ester of a polyvalent alcohol with at least one fatty acid chosen from a group consisting of caprylic esters of propylene glycol;

said surfactant and co-surfactant being in a the ratio by weight TA/CoTA being between 0.2 and 6.[[,]]

characterized in that the ester of a polyvalent alcohol with at least one fatty acid in the cosurfactant phase is chosen from the group comprising caprylic esters of propylene glycol.

15. (Currently amended) Composition according to Claim 14, characterized in that the wherein said lipophilic phase comprises a mixture of glycerol mono, di- and triesters and of PEG mono- and diesters with the combination of saturated C₈-C₁₈ fatty acids, the said mixture having has an HLB value equal to 14 and it represents representing between 50 and 95% by weight of the composition.

- 16. (Currently amended) A composition Composition according to claim 14, characterized in that wherein the surfactant phase represents between 1% and 30% by weight of the mixture composition.
- 17. (Currently amended) <u>A composition Composition</u> according to claim 14, characterized in that wherein the co-surfactant phase represents between 3% and 32% by weight of the mixture composition.
- 18. (Currently amended) <u>A composition</u> Composition according to claim 14, characterized in that the <u>wherein said</u> active principle <u>is a belongs to the</u> statin family.
- 19. (Currently amended) <u>A composition</u> according to Claim 18, characterized in that the wherein said statin is simvastatin.
- 20. (Currently amended) A composition Composition according to Claim 19, characterized in that the wherein said simvastatin represents between 0.1% and 6% by weight of the composition and advantageously 4% by weight.
- 21. (Currently amended) A composition Composition according to claim 14, characterized in that it comprises by weight wherein said active principle is simvastatin and wherein said simvastatin represents by weight between 0.1% and 6% of the composition, said lipophilic phase represents by weight between 52% and 70% of the composition, said surfactant phase represents by weight between 5% and 30% of the composition, and said co-surfactant phase is comprised of propylene glycol monocaprylate, wherein propylene glycol monocaprylate represents by weight between 15% and 30% of the composition. ÷

- between 0.1% and 6% of simvastatin,

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between 52% and 70% of Gélucire® 44/14,
- between 5% and 30% of Labrasol®,
between 15% and 30% of propylene glycol monocaprylate.
22. (Currently amended) A composition Composition according to Claim 21, characterized in
that the wherein said propylene glycol monocaprylate consists of Capryol® PGMC representing
represents by weight between 15% and 25% by weight of the composition.
23. (Currently amended) A composition Composition according to Claim 21, characterized in that the wherein said propylene glycol monocaprylate consists of Capryol® 90 representing
represents by weight between 20% and 30% by weight of the composition.
24. (Currently amended) <u>A composition Composition</u> according to Claim 21, characterized in that the wherein said surfactant and co-surfactant are in a 0.5 ratio by weight TA/CoTA is equal to 0,5.
25. (New) A composition according to claim 21, wherein said lipophilic phase has an HLB value of 14.
26. (New) A composition according to claim 21, wherein said lipophilic phase is comprising of lauric macrogolglycerides.
27. (New) A composition according to claim 21, wherein said surfactant phase has an HLB value between 5 and 20.
28. (New) A composition according to claim 21, wherein said surfactant phase is caprylocapric

magrogol glyceride.